Doc code: RCEX

Doc description: Request for Continued Examination (RCE)

submission even if this box is not checked.

Patent Practitioner Signature

Applicant Signature

PTO/SB/30EFS (03/08)
Approved for use through 03/31/2008 OMB 0651-0631
U.S. Patent and Trademark Office; U.S. DEPARTMENT OF COMMERCE

U.S. Patent and Trademark Office; U.S. DEPARTMENT OF COMMERCE
Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it contains a valid OMB control number.

REQUEST FOR CONTINUED EXAMINATION(RCE)TRANSMITTAL (Submitted Only via EFS-Web)

(Submitted Siny via Li S-vieb)								
Application	10766964	Filing	2004-01-29	Docket Number	E03.001/U	Art	2176	
Number	10700904	Date	2004-01-29	(if applicable)	E03.001/0	Unit	2170	
First Named William A. Margiloff				Examiner	Henry W. Orr			
Inventor William A. Marginum		Name						

This is a Request for Continued Examination (RCE) under 37 CFR 1.114 of the above-identified application.

Consider the arguments in the Appeal Brief or Reply Brief previously filed on

Request for Continued Examination (RCE) practice under 37 CFR 1.114 does not apply to any utilify or plant application filed prior to June 8, 1995, or to any design application. The Instruction Sheet for this form is located at WWW.USPTO.GOV

SUBMISSION REQUIRED UNDER 37 CFR 1.114 Note: If the RCE is proper, any previously filed unentered amendments and amendments enclosed with the RCE will be entered in the order

in which they wers filled unless applicant instructs otherwise. If applicant does not wish to have any previously filed unentered amendment(s) entered, applicant must request non-entry of such amendment(s).

— Previously submitted. If a final Office action is outstanding, any amendments filed after the final Office action may be considered as a

	Other					
\boxtimes	Enclosed					
	☐ Information Disclosure Statement (IDS)					
	Affidavit(s)/ Declaration(s)					
	○ Other Petition for 3-Month Extension of Time					
	MISCELLANEOUS					
	Suspension of action on the above-identified application is requested under 37 CFR 1.103(c) for a period of months (Period of suspension shall not exceed 3 months; Fee under 37 CFR 1.17(i) required)					
	Other					
	FEES					
⊠	The RCE fee under 37 CFR 1.17(e) is required by 37 CFR 1.114 when the RCE is filled. The Director is hereby authorized to charge any underpayment of fees, or credit any overpayments, to Deposit Account No 501852					
	SIGNATURE OF APPLICANT, ATTORNEY, OR AGENT REQUIRED					

Doc description: Request for Continued Examination (RCE)

PTO/SB/30EFS (03/08)
Approved for use through 03/31/2008 OMB 0651-0031
U.S. Patent and Trademark Office; U.S. DEPARTMENT OF COMMERCE

Under the Paperwork Reduction Act of 1995, no persons are required to respond to a control or of the Paperwork Reduction Act of 1995, no persons are required to respond to a control or of the Paperwork Reduction Act of 1995, no persons are required to respond to a control or of the Paperwork Reduction Act of 1995, no persons are required to respond to a control or of the Paperwork Reduction Act of 1995, no persons are required to respond to a control or of the Paperwork Reduction Act of 1995, no persons are required to respond to a control or of the Paperwork Reduction Act of 1995, no persons are required to respond to a control or of the Paperwork Reduction Act of 1995, no persons are required to respond to a control or of the Paperwork Reduction Act of 1995, no persons are required to respond to a control or of the Paperwork Reduction Act of 1995, no persons are required to respond to a control or of the Paperwork Reduction Act of 1995, no persons are required to respond to a control or of the Paperwork Reduction Act of 1995, no persons are required to respond to a control or of the Paperwork Reduction Act of 1995, no persons are required to respond to a control or of the Paperwork Reduction Act of 1995, no persons are required to respond to a control or of the Paperwork Reduction Act of 1995, no persons are required to respond to the Paperwork Reduction Act of 1995, no persons are required to respond to the Paperwork Reduction Act of 1995, no persons are required to respond to the Paperwork Reduction Act of 1995, no persons are required to respond to the Paperwork Reduction Act of 1995, no persons are required to respond to the Paperwork Reduction Act of 1995, no persons are required to respond to the Paperwork Reduction Act of 1995, no persons are required to respond to the Paperwork Reduction Act of 1995, no persons are required to the Paperwork Reduction Act of 1995, no persons are required to the Paperwork Reduction Act of 1995, no persons are required to the Paperwork Reduction Act of 1995, no persons a

Signature of Registered U.S. Patent Practitioner							
Signature	/Patrick J. Buckley/	Date (YYYY-MM-DD)	2008-03-05				
Name	Patrick J. Buckley	Registration Number	40928				

This collection of information is required by 37 CFR 1.114. The information is required to obtain or retain a benefit by the public which is to fine (and by the USPTO to process) an application. Confidentiality is owerened by 35 U S C. 122 and 37 CFR 1.11 and 1.14. This collection is estimated to take 12 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450.

If you need assistance in completing the form, call 1-800-PTO-9199 and select option 2.

Privacy Act Statement

The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the attached form related to a patient application or patient. Accordingly, pursuant to the requirements of he Act, please be advised that: (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) furnishing of the information solicited is voluntary; and (3) the principal purpose for which the information is used by the U.S. Patient and Trademark Office is to process ana/for examine your submission related to a patient application or patient. If you do not furnish the requested information, the U.S. Patient and Trademark Office may not be able to process and/or examine your submission, which may result in termination of proceedings or abandomment of the application or expiration of the patient.

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information
 Act (6 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the
 Department of Justice to determine whether the Freedom of Information Act requires disclosure of these records.
- A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement necodiations.
- A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record pertains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
- A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need
 for the information in order to perform a contract. Recipients of information shall be required to comply with the
 requirements of the Privacy Act of 1974, as amended, pursuant to 5 U.S.C. 552a(m).
- A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
 may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization,
 pursuant to the Patent Cooperation Treaty.
- A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
- 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or his/her designee, during an inspection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 44 U.S.C. 2904 and 2906. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
- 8. A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S.C. 12(2b) or issuance of a patent pursuant to 35 U.S.C. 151. Further, a record may be disclosed, subject to the limitations of 37 CFR 1.14, as a routine use, to the public if the record was filed in an application which became abandoned or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inspections or an issued patent.
- A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.